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AMENDMENTS TO THE CLAIMS

Listing of Claims:

1. (Currently amended) A method comprising:

a) providing one or more coded oligonucleotide probes, each coded oligonucleotide

probe comprising an oligonucleotide attached to at least one unique nanocode wherein each

nanocode comprises a detectable non-encoding feature, which detectable non-encoding feature

comprises a feature tag pattern, and wherein the tag pattern provides a quality control check for

detecting nanocodes and/or distinguishes target nucleotides from self-assembled coded

oligonucleotide probe structures;

contacting at least one target nucleic acid with the one or more coded oligonucleotide b)

probes; and

c) utilizing the feature tag to provide a quality control check for detecting nanocodes

and/or distinguishes target nucleotides from self-assembled coded oligonucleotide probe structures;

and

identifying coded oligonucleotide probes that bind to the target nucleic acid using

scanning probe microscopy (SPM) to detect the nanocode and the detectable non-encoding feature

tag.

2. (Currently amended) The method of claim 1, wherein the one or more coded

oligonucleotide probes comprise permutations of a linear order of nucleic acid residues, which

linear order substantially represents all possible complementary sequences for a particular length of

oligonucleotide.

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- 3. (Original) The method of claim 1, wherein the nanocode is selected from the group consisting of carbon nanotubes, fullerenes, submicrometer metallic barcodes, nanoparticles and quantum dots.
- 4. (Original) The method of claim 1, wherein the nucleic acid is attached to a surface.
- 5. (Original) The method of claim 4, further comprising ligating adjacent coded probes that are hybridized to the nucleic acid.
- 6. (Previously Presented) The method of claim 5, further comprising separating ligated coded probes from the target nucleic acid and non-ligated coded probes.
- 7. (Original) The method of claim 6, wherein the ligated coded probes form reading frames.
- 8. (Original) The method of claim 1, further comprising aligning the coded probes on a surface by molecular combing.
- 9. (Previously Presented) The method of claim 1, wherein the scanning probe microscopy is atomic force microscopy, scanning tunneling microscopy, lateral force microscopy,

chemical force microscopy, force modulation imaging, magnetic force microscopy, high frequency magnetic force microscopy, magnetoresistive sensitivity mapping, electric force microscopy, scanning capacitance microscopy, scanning spreading resistance microscopy, tunneling atomic force microscopy or conductive atomic force microscopy.

- 10. (Previously Presented) The method of claim 2, further comprising determining the nucleotide sequences of oligonucleotides that bind to the target nucleic acid.
- 11. (Previously Presented) The method of claim 10, further comprising determining a nucleotide sequence of the target nucleic acid from the sequences of oligonucleotides that bind to the target nucleic acid.
- 12. (Previously Presented) The method of claim 1, further comprising identifying the target nucleic acid from the coded probes that bind to the target nucleic acid.
- 13. (Original) The method of claim 1, wherein two or more target nucleic acids are present in a sample.
- 14. (Previously Presented) The method of claim 1, wherein at least two target nucleic acids are contacted in the sample at the same time.

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15. (Currently amended) The method of claim 1, wherein the detectable non-

encoding feature tag is provided by a detectable feature tag associated with the nanocode.

16. (Currently amended) The method of claim 15 wherein the detectable non-encoding

feature tag comprises a start tag.

17. (Original) The method of claim 1, further comprising transforming the molecular

nanocode to form a decompressed nanocode.

18. (Currently amended) The method of claim 1, wherein the detectable feature tag is

comprises a checksum barcode segment.

19. (Currently amended) The method of claim 1, wherein the detectable feature tag

comprises a header segment and an encoding segment.

20. (Currently amended) A composition comprising at least one coded probe, each

coded probe comprising a probe molecule attached to at least one nanocode comprising a detectable

non-encoding feature, which detectable non-encoding feature comprises a feature tag pattern,

wherein the feature tag pattern provides has a property to provide a quality control check for

detecting nanocodes and/or distinguishes target nucleotides from self-assembled coded

oligonucleotide probe structures, the nanocode being detectable using a single molecule level

surface analysis method.

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21. (Previously Presented) The composition of claim 20, wherein the probe molecule is an oligonucleotide, a polynucleotide, a nucleic acid, an antibody, an antibody fragment, a genetically engineered antibody, a single chain antibody, a humanized antibody, a protein, a receptor, a transcription factor, a peptide, a lectin, a substrate, an inhibitor, an activator, a ligand, a hormone, a cytokine, a chemokine, or a pharmaceutical.

- 22. (Original) The composition of claim 20, wherein the probe molecule is an oligonucleotide.
- 23. (Original) The composition of claim 20, wherein the nanocode is selected from the group consisting of carbon nanotubes, fullerenes, submicrometer metallic barcodes, nanoparticles and quantum dots.
- 24. (Currently amended) The composition of claim 20, wherein the detectable non-encoding feature tag is comprises a start tag.
- 25. (Original) The composition of claim 20, wherein the nanocode is a compressed nanocode.
- 26. (Original) The composition of claim 20, wherein the nanocode comprises reading frames.

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- 27. (Original) The composition of claim 20, wherein the nanocode comprises a header region and an encoding region.
- 28. (Original) The composition of claim 20, wherein the nanocode is detectable using scanning probe microscopy (SPM).
 - 29. (Currently amended) A system comprising:
 - a) a scanning probe microscope (SPM);
 - b) a surface; and
- c) at least one coded oligonucleotide probe attached to the surface, wherein the coded oligonucleotide probe comprises a nanocode comprising a detectable non-encoding feature, which detectable non-encoding feature comprises a feature tag pattern, and wherein the feature tag pattern provides has a property to provide a quality control check for detecting nanocodes and/or distinguishes target nucleotides from self-assembled coded oligonucleotide probe structures, the nanocode being detectable using SPM.
- 30. (Original) The system of claim 29, wherein the coded oligonucleotide probes comprise ligated oligonucleotides.
- 31. (Original) The system of claim 30, wherein the ligated oligonucleotides form reading frames.

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- 32. (Original) The system of claim 29, wherein the scanning probe microscope is an atomic force microscope or a scanning tunneling microscope.
- 33. (Currently amended) The system of claim 29, wherein the detectable non-encoding feature tag is comprises a start tag.
- 34. (Currently amended) The system of claim 29, wherein the nanocode is comprises a compressed nanocode.
- 35. (Original) The system of claim 29, wherein the nanocode comprises reading frames.
- 36. (Original) The system of claim 29, wherein the nanocode comprises a header region and an encoding region.